

# Sr. Project Manager, Engineering

Job ID  
REQ-10076496  
апр 29, 2026  
США

## Сводка

Bring your experience, leadership, and strategic mindset to a role where your work directly enables the delivery of innovative therapies to patients around the world. As a Sr. Project Manager, Engineering at Novartis, you will lead a complex portfolio of capital projects within a regulated manufacturing environment, guiding teams through ambiguity and technical challenges with confidence. You will serve as a trusted partner to site leadership, drive decisions that shape long-term manufacturing strategy, and champion operational excellence while balancing cost, quality, and schedule. This role offers the opportunity to influence at scale, mentor teams, and leave a lasting footprint across the site's capital portfolio.

## About the Role

### Key Responsibilities:

- Lead multiple capital projects from initiation through close-out, ensuring delivery on time, within budget, and to quality standards
- Act as primary client contact, proactively managing stakeholder expectations and maintaining strong working relationships
- Manage project scope, schedule, and budget changes with clear impact assessments and documented approvals
- Oversee contractors and internal teams, ensuring safe, compliant, and effective project execution
- Resolve project issues, risks, and resource constraints in partnership with site leadership
- Prepare and manage project budgets, forecasts, and funding documentation
- Develop project objectives aligned with business plans and user requirements
- Coordinate planning and scheduling across project teams and manufacturing operations
- Communicate project performance, risks, and mitigation plans to site and senior leadership
- Drive operational excellence by capturing lessons learned and supporting continuous improvement initiatives

### Essential Requirements:

- Bachelor's degree in Chemical, Electrical, or Mechanical Engineering, or a related technical field, or equivalent experience
- Nine years of experience supporting pharmaceutical or biopharmaceutical manufacturing operations in regulated environments
- Demonstrated experience leading FDA-regulated and GMP-compliant projects within highly regulated facilities
- Strong knowledge of FDA regulations and Good Manufacturing Practice systems
- Experience delivering capital improvement projects, including process and production layout development
- Experience managing Good Manufacturing Practice construction activities in brownfield and renovation environments
- Applied knowledge of Quality by Design, Six Sigma, and operational excellence methodologies
- Excellent written and verbal communication skills, including strong technical writing capabilities

### Novartis Compensation and Benefit Summary:

The salary for this position is expected to range between \$108,500 and \$201,500 annually

The final salary offered is determined based on factors like, but not limited to, relevant skills and experience, and upon joining Novartis will be reviewed periodically. Novartis may change the published salary range based on company and market factors.

Your compensation will include a performance-based cash incentive and, depending on the level of the role, eligibility to be considered for annual equity awards.

US-based eligible employees will receive a comprehensive benefits package that includes health, life and disability benefits, a 401(k) with company contribution and match, and a variety of other benefits. In addition, employees are eligible for a generous time off package including vacation, personal days, holidays and other leaves.

#LI-Onsite

**Why Novartis:** Helping people with disease and their families takes more than innovative science. It takes a community of smart, passionate people like you. Collaborating, supporting and inspiring each other. Combining to achieve breakthroughs that change patients' lives. Ready to create a brighter future together?  
<https://www.novartis.com/about/strategy/people-and-culture>

**Benefits and Rewards:** Learn about all the ways we'll help you thrive personally and professionally.

[Read our handbook \(PDF 30 MB\)](#)

### EEO Statement:

The Novartis Group of Companies are Equal Opportunity Employers. We do not discriminate in recruitment, hiring, training, promotion or other employment practices for reasons of race, color, religion, sex, national origin, age, sexual orientation, gender identity or expression, marital or veteran status, disability, or any other legally protected status.

### Accessibility & Reasonable Accommodations

The Novartis Group of Companies are committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the application process, or to perform the essential functions of a position, please send an e-mail to [us.reasonableaccommodations@novartis.com](mailto:us.reasonableaccommodations@novartis.com) or call +1(877)395-2339 and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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Production / Manufacturing  
Место  
США  
Состояние  
North Carolina  
Сайт  
Durham  
Company / Legal Entity  
U473 (FCRS = US473) Novartis Gene Therapies  
Functional Area  
Technical Operations  
Job Type  
Full time  
Employment Type  
Regular  
Shift Work  
No

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